

**AZEOTROPE-LIKE COMPOSITIONS OF  
TETRAFLUOROPROPENE AND TRIFLUOROIODOMETHANE****CROSS-REFERENCES TO RELATED APPLICATIONS**

5 This application claims the benefit of priority to the U.S. applications identified by Attorney Docket Nos. H0006721-4510, H0005517-4510, H0005706-4510, H0007522-4510, and H0007523-4510, respectively, all of which were filed on even date herewith and are incorporated herein by reference.

**FIELD OF INVENTION**

10 The present invention provides azeotrope-like compositions of *trans*-1,1,1,3-tetrafluoropropene and trifluoroiodomethane, and uses thereof.

**BACKGROUND**

15 Fluorocarbon based fluids have found widespread use in industry in a number of applications, including as refrigerants, aerosol propellants, blowing agents, heat transfer media, and gaseous dielectrics. Because of the suspected environmental problems associated with the use of some of these fluids, including the relatively high global warming potentials associated therewith, it is desirable to use fluids having low or even zero ozone depletion  
20 potential, such as hydrofluorocarbons ("HFCs"). Thus, the use of fluids that do not contain chlorofluorocarbons ("CFCs") or hydrochlorofluorocarbons ("HCFCs") is desirable. Additionally, the use of single component fluids or azeotropic mixtures, which do not fractionate on boiling and evaporation, is desirable. However, the identification of new, environmentally-safe, non-fractionating mixtures is complicated due to the fact that azeotrope  
25 formation is not readily predictable.

The industry is continually seeking new fluorocarbon based mixtures that offer alternatives, and are considered environmentally safer substitutes for CFCs and HCFCs. Of particular interest are mixtures containing both hydrofluorocarbons and other fluorinated compounds, both of low ozone depletion potentials. Such mixtures are the subject of this  
30 invention.

**DESCRIPTION OF PREFERRED EMBODIMENTS**

The present inventors have developed several compositions that help to satisfy the continuing need for alternatives to CFCs and HCFCs. According to certain embodiments, the

present invention provides azeotrope-like compositions comprising *trans*-1,1,1,3-tetrafluoropropene (“HFO-1234ze”) and trifluoroiodomethane (“CF<sub>3</sub>I”).

The preferred compositions of the invention tend both to be non-flammable and to exhibit relatively low global warming potentials (“GWPs”). Accordingly, applicants have recognized that such compositions can be used to great advantage in a number of applications, including as replacements for CFCs, HCFCs, and HFCs (such as HFC-134a) in refrigerant, aerosol, and other applications.

Additionally, applicants have recognized surprisingly that azeotrope-like compositions of HFO-1234ze and CF<sub>3</sub>I can be formed. Accordingly, in other embodiments, the present invention provides methods of producing an azeotrope-like composition comprising combining HFO-1234ze and CF<sub>3</sub>I in amounts effective to produce an azeotrope-like composition.

In addition, applicants have recognized that the azeotrope-like compositions of the present invention exhibits properties that make that make them advantageous for use as, or in, refrigerant compositions. Accordingly, in yet other embodiments, the present invention provides refrigerant compositions comprising an azeotrope-like composition of HFO-1234ze and CF<sub>3</sub>I.

#### Azeotrope-like Compositions

As used herein, the term “azeotrope-like” is intended in its broad sense to include both compositions that are strictly azeotropic and compositions that behave like azeotropic mixtures. From fundamental principles, the thermodynamic state of a fluid is defined by pressure, temperature, liquid composition, and vapor composition. An azeotropic mixture is a system of two or more components in which the liquid composition and vapor composition are equal at the stated pressure and temperature. In practice, this means that the components of an azeotropic mixture are constant-boiling and cannot be separated during a phase change.

The azeotrope-like compositions of the invention may include additional components that do not form new azeotrope-like systems, or additional components that are not in the first distillation cut. The first distillation cut is the first cut taken after the distillation column displays steady state operation under total reflux conditions. One way to determine whether the addition of a component forms a new azeotrope-like system so as to be outside of this invention is to distill a sample of the composition with the component under conditions that would be expected to separate a non-azeotropic mixture into its separate components. If the mixture containing the additional component is non-azeotrope-like, the additional component

will fractionate from the azeotrope-like components. If the mixture is azeotrope-like, some finite amount of a first distillation cut will be obtained that contains all of the mixture components that is constant boiling or behaves as a single substance.

It follows from this that another characteristic of azeotrope-like compositions is that there is a range of compositions containing the same components in varying proportions that are azeotrope-like or constant boiling. All such compositions are intended to be covered by the terms "azeotrope-like" and "constant boiling". As an example, it is well known that at differing pressures, the composition of a given azeotrope will vary at least slightly, as does the boiling point of the composition. Thus, an azeotrope of A and B represents a unique type of relationship, but with a variable composition depending on temperature and/or pressure. It follows that, for azeotrope-like compositions, there is a range of compositions containing the same components in varying proportions that are azeotrope-like. All such compositions are intended to be covered by the term azeotrope-like as used herein.

It is well-recognized in the art that it is not possible to predict the formation of azeotropes. (See, for example, U.S. Patent No. 5,648,017 (column 3, lines 64-65) and U.S. Patent No. 5,182,040 (column 3, lines 62-63), both of which are incorporated herein by reference). Applicants have discovered unexpectedly that HFO-1234ze and CF<sub>3</sub>I form azeotrope-like compositions.

According to certain preferred embodiments, the azeotrope-like compositions of the present invention comprise, and preferably consist essentially of, effective azeotrope-like amounts of HFO-1234ze and CF<sub>3</sub>I. The term "effective azeotrope-like amounts" as used herein refers to the amount of each component which upon combination with the other component, results in the formation of an azeotrope-like composition of the present invention. Preferably, the present azeotrope-like compositions comprise, and preferably consist essentially of, from greater than zero to about 85 weight percent HFO-1234ze and from about 15 to less than 100 weight percent of CF<sub>3</sub>I. More preferably, the azeotrope-like compositions comprise, and preferably consist essentially of, from greater than zero to about 80 weight percent HFO-1234ze and from about 20 to less than 100 weight percent of CF<sub>3</sub>I, more preferably from about 1 to about 40 weight percent HFO-1234ze and from about 60 to about 99 weight percent of CF<sub>3</sub>I, even more preferably from about 5 to about 35 weight percent HFO-1234ze and from about 65 to about 95 weight percent of CF<sub>3</sub>I, and even more preferably from about 15 to about 25 weight percent HFO-1234ze and from about 85 to about 75 weight percent of CF<sub>3</sub>I. Unless otherwise indicated, the weight percents disclosed herein are based on the total weight of CF<sub>3</sub>I, and HFO-1234ze in a composition.

The azeotrope-like compositions described herein preferably have a boiling point of from about -25°C to about -21°C at a pressure of about 14.42 psia. In certain more preferred embodiments, the present azeotrope-like compositions have a boiling point of from about -25°C to about -22°C at a pressure of about 14.42 psia, and in even more preferred  
5 embodiments, from about -24°C to about -22°C at a pressure of about 14.42 psia.

The azeotrope-like compositions of the present invention can be produced by combining effective azeotrope-like amounts of HFO-1234ze and CF<sub>3</sub>I. Any of a wide variety of methods known in the art for combining two or more components to form a composition can be adapted for use in the present methods to produce an azeotrope-like composition. For  
10 example, HFO-1234ze and CF<sub>3</sub>I can be mixed, blended, or otherwise contacted by hand and/or by machine, as part of a batch or continuous reaction and/or process, or via combinations of two or more such steps. In light of the disclosure herein, those of skill in the art will be readily able to prepare azeotrope-like compositions according to the present invention without undue experimentation.

#### Composition Additives

The azeotrope-like compositions of the present invention may further include any of a variety of optional additives including stabilizers, metal passivators, corrosion inhibitors, and the like.

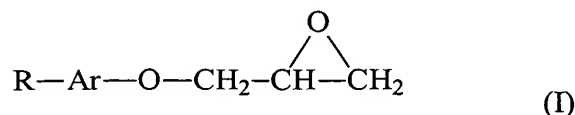
According to certain embodiments, the azeotrope-like compositions of the present invention further comprise a stabilizer. Any of a variety of compounds suitable for stabilizing an azeotrope-like composition of the present invention may be used. Examples of certain preferred stabilizers include stabilizer compositions comprising at least one phenol composition and at least one epoxide selected from the group consisting of aromatic  
25 epoxides, alkyl epoxides, alkenyl epoxides, and combinations of two or more thereof.

Any of a variety of phenol compounds are suitable for use in the present compositions. While applicants do not wish to be bound by or to any theory of operation, it is believed that the present phenols act as radical scavengers in the CF<sub>3</sub>I compositions and thereby tend to increase the stability of such compositions. As used herein the term “phenol compound” refers generally to any substituted or unsubstituted phenol. Examples of suitable  
30 phenol compounds include phenols comprising one or more substituted or unsubstituted cyclic, straight-chain, or branched aliphatic substituent group, such as, alkylated monophenols including: 2,6-di-tert-butyl-4-methylphenol; 2,6-di-tert-butyl-4-ethylphenol;

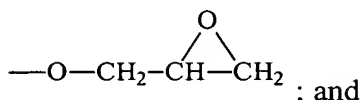
2,4-dimethyl-6-tert-butylphenol; tocopherol; and the like, hydroquinone and alkylated hydroquinones including: t-butyl hydroquinone; other derivatives of hydroquinone; and the like, hydroxylated thiodiphenyl ethers including: 4,4'-thiobis (2-methyl-6-tert-butylphenol); 4,4'-thiobis (3-methyl-6-tert-butylphenol); 2,2'-thiobis (4-methyl-6-tert-butylphenol); and the like, alkylidene-bisphenols including: 4,4'-methylenebis(2,6-di-tert-butylphenol); 4,4'-bis(2,6-di-tert-butylphenol); derivatives of 2,2- or 4,4-biphenyldiols; 2,2'-methylenebis(4-ethyl-6-tert-butylphenol); 2,2'-methylenebis(4-methyl-6-tert-butylphenol); 4,4'-butylidenebis(3-methyl-6-tert-butylphenol); 4,4'-isopropylidenebis(2,6-di-tert-butylphenol); 2,2'-methylenebis(4-methyl-6-nonylphenol); 2,2'-isobutylidenebis(4,6-dimethylphenol); 2,2'-methylenebis(4-methyl-6-cyclohexylphenol), 2,2- or 4,4-biphenyldiols including 2,2'-methylenebis(4-ethyl-6-tert-butylphenol), butylated hydroxy toluene (BHT), bisphenols comprising heteroatoms including: 2,6-di-tert-butyl-4-dimethylamino-p-cresol; 4,4'-thiobis(6-tert-butyl-m-cresol); and the like; acylaminophenols; 2,6-di-tert-butyl-4(N,N'-dimethylaminomethylphenol); sulfides including: bis(3-methyl-4-hydroxy-5-tert-butylbenzyl)sulfide; bis(3,5-di-tert-butyl-4-hydroxybenzyl)sulfide; and the like; as well as, phenolic UV absorb and light stabilizers. Certain preferred phenols include alkylated monophenols such as tocopherol, BHT, hydroquinones, and the like. Certain particularly preferred phenols include tocopherol, and the like. Most phenols are commercially available. A single phenol compound and/or mixtures of two or more phenols may be used in the present compositions.

Any of a variety of epoxides are suitable for use in the compositions of the present invention. While applicants do not wish to be bound by or to any theory of operation, it is believed that the epoxides of the present invention act as acid scavengers in the CF<sub>3</sub>I compositions and thereby tend to increase the stability of such compositions. A single aromatic epoxide and/or mixtures of two or more aromatic epoxides may be used in the present compositions.

Examples of suitable aromatic epoxides include those defined by the formula I below:



wherein: R is hydrogen, hydroxyl, alkyl, fluoroalkyl, aryl, fluoroaryl, or



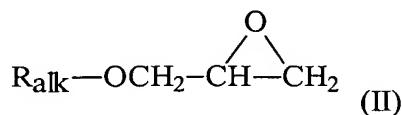
Ar is a substituted or unsubstituted phenylene or naphthylene moiety. Certain preferred aromatic epoxides of Formula I include those wherein Ar is phenylene or phenylene substituted with one or more substituents including alkyls, alkenyls, alkynyls, aryls, alkylaryls, halogens, halogenated alkyls, halogenated alkenyls, halogenated alkynyls, halogenated aryls, halogenated arylalkyls, hydroxyls, heteroatom moieties, and the like. Examples of suitable compounds of Formula I wherein Ar is an unsubstituted or substituted phenylene include butylphenylglycidyl ether; pentylphenylglycidyl ether; hexylphenylglycidyl ether; heptylphenylglycidyl ether; octylphenylglycidyl ether; nonylphenylglycidyl ether; decylphenylglycidyl ether; glycidyl methyl phenyl ether; 1,4-diglycidyl phenyl diether; 4-methoxyphenyl glycidyl ether; derivatives thereof; and the like.

Certain other preferred aromatic epoxides of Formula I include those wherein Ar is naphthylene or naphthylene substituted with one or more substituents including alkyls, alkenyls, alkynyls, aryls, alkylaryls, halogens, halogenated alkyls, halogenated alkenyls, halogenated alkynyls, halogenated aryls, halogenated arylalkyls, hydroxyls, heteroatom moieties, and the like. Examples of suitable compounds of Formula I wherein Ar is an unsubstituted or substituted naphthylene include naphthyl glycidyl ether; 1,4-diglycidyl naphthyl diether; derivatives thereof; and the like.

Examples of other suitable aromatic epoxides include bisoxiranes, such as, 2,2'[[[5-heptadecafluorooctyl]1,3phenylene]bis[[2,2,2trifluoromethyl]ethylidene]oxymethylene] bisoxirane; and the like.

In certain preferred embodiments, the aromatic epoxides for use in the present invention comprise an epoxide of Formula I wherein Ar is phenylene, substituted phenylene, naphthylene, or substituted naphthylene. More preferably, the aromatic epoxides comprise an epoxide of Formula I wherein Ar is phenylene or substituted phenylene. Examples of certain more preferred aromatic epoxides include butylphenyl glycidyl ether, and the like.

Any of a variety of alkyl and/or alkenyl epoxides are suitable for use in the present compositions. Examples of suitable alkyl and alkenyl epoxides include those of Formula II:



wherein  $\text{R}_{\text{alk}}$  is a substituted or unsubstituted alkyl or alkenyl group. Certain preferred epoxides of Formula II comprise alkyl epoxide compounds wherein  $\text{R}_{\text{alk}}$  is an alkyl group having from about 1 to about 10 carbon atoms, more preferably from about 1 to about 6 carbon atoms, and wherein the alkyl may be unsubstituted or further substituted with one or

more substituents including alkyls, alkenyls, alkynyls, aryls, alkylaryls, halogens, halogenated alkyls, halogenated alkenyls, halogenated alkynyls, halogenated aryls, halogenated arylalkyls, hydroxyls, heteroatom moieties, and the like. Examples of such preferred alkyl epoxides of Formula II include n-butyl glycidyl ether, isobutyl glycidyl ether, 5 hexanediol diglycidyl ether, and the like, as well as, fluorinated and perfluorinated alkyl epoxides, and the like. Certain more preferred alkyl epoxides comprise hexanediol diglycidyl ether, and the like.

Certain other preferred epoxides of Formula II comprise alkenyl epoxide compounds wherein  $R_{alk}$  is an alkenyl group having from about 1 to about 10 carbon atoms, more 10 preferably from about 1 to about 6 carbon atoms, and wherein the alkenyl may be unsubstituted or further substituted with one or more substituents including alkyls, alkenyls, alkynyls, aryls, alkylaryls, halogens, halogenated alkyls, halogenated alkenyls, halogenated alkynyls, halogenated aryls, halogenated arylalkyls, hydroxyls, heteroatom moieties, and the like. Examples of such preferred alkenyl epoxides of Formula II include allyl glycidyl ether, 15 fluorinated and perfluorinated alkenyl epoxides, and the like. More preferred alkenyl epoxides include allyl glycidyl ether, and the like. A single alkyl epoxide or alkenyl epoxide and/or combinations of two or more thereof may be used in the present compositions.

In certain other preferred embodiments, the alkyl epoxide for use as an acid scavenger in the present composition comprises polypropylene glycol diglycidyl ether. Examples of 20 polypropylene glycol diglycidyl ether suitable for use in the present invention includes the ether available commercially from SACHEM, Europe.

In addition, in certain embodiments, the epoxide for use in the present invention comprises combinations of two or more aromatic, alkyl, and/or alkenyl substituents. Such epoxides are referred to generally as "multisubstituted epoxides."

25 According to certain preferred embodiments, the stabilizer for use in the present invention comprises a combination of at least one phenol compound and at least one aromatic, alkyl, or alkenyl epoxide. Examples of suitable combinations include stabilizers comprising: tocopherol and allyl glycidyl ether, BHT and glycidyl butyl ether, and the like. Certain particularly preferred combinations include stabilizers comprising: tocopherol and 30 allyl glycidyl ether, and the like.

Any suitable relative amount of the at least one phenol compound and the at least one aromatic, alkyl, or alkenyl epoxide may be used in the preferred stabilizers. For example, the weight ratio of phenol compound(s) to aromatic or fluorinated alkyl epoxide(s) can be varied from about 1:99 to about 99:1. In certain preferred embodiments, the weight ratios of phenol

compound(s) to aromatic, alkyl, alkenyl, multisubstituted, or fluorinated alkyl epoxide(s) is from about 30 to about 1, more preferably from about 7 to about 1, more preferably from about 2 to about 1, and even more preferably about 1:1.

Any suitable effective amount of stabilizer may be used in the trifluoroiodomethane compositions of the present invention. As used herein, the term "effective stabilizing amount" refers to an amount of stabilizer of the present invention which, when added to a composition comprising trifluoroiodomethane, results in a stabilized composition wherein the trifluoroiodomethane therein degrades more slowly and/or to a lesser degree relative to the original composition, under the same, or similar, conditions. In certain preferred embodiments, an "effective stabilizing amount" of stabilizer comprises an amount which, when added to a composition comprising trifluoroiodomethane, results in a stabilized composition wherein the trifluoroiodomethane therein degrades more slowly and/or to a lesser degree relative to the original composition under the conditions of at least one, or both, of the standards tests SAE J1662 (issued June 1993) and/or ASHRAE 97-1983R. In certain more preferred embodiments, an "effective stabilizing amount" of stabilizer comprises an amount which, when added to a composition comprising trifluoroiodomethane, results in a composition having a stability that is at least as good as, if not better, than the stability of a comparable composition comprising dichlorodifluoromethane (R-12) in mineral oil, under at least one of the standard tests SAE J1662 (issued June 1993) and/or ASHRAE 97-1983R. Certain preferred effective amounts of stabilizer for use in the present invention comprise from about 0.001 to about 10, more preferably from about 0.01 to about 5, even more preferably from about 0.3 to about 4 weight percent, and even more preferably from about 0.3 to about 1 weight percent based on the total weight of trifluoroiodomethane in the composition of the present invention.

In certain preferred embodiments, the compositions of the present invention further comprise a lubricant. Any of a variety of conventional lubricants may be used in the compositions of the present invention. An important requirement for the lubricant is that, when in use in a refrigerant system, there must be sufficient lubricant returning to the compressor of the system such that the compressor is lubricated. Thus, suitability of a lubricant for any given system is determined partly by the refrigerant/lubricant characteristics and partly by the characteristics of the system in which it is intended to be used. Examples of suitable lubricants include mineral oil, alkyl benzenes, polyol esters, including polyalkylene glycols, PAG oil, and the like. Mineral oil, which comprises paraffin oil or naphthenic oil, is commercially available. Commercially available mineral oils include Witco LP 250



(registered trademark) from Witco, Zerol 300 (registered trademark) from Shrieve Chemical, Sunisco 3GS from Witco, and Calumet R015 from Calumet. Commercially available alkyl benzene lubricants include Zerol 150 (registered trademark). Commercially available esters include neopentyl glycol dipelargonate which is available as Emery 2917 (registered trademark) and Hatcol 2370 (registered trademark). Other useful esters include phosphate esters, dibasic acid esters, and fluoroesters. Preferred lubricants include polyalkylene glycols and esters. Certain more preferred lubricants include polyalkylene glycols.

#### Uses of the Compositions

The present compositions have utility in a wide range of applications. For example, one embodiment of the present invention relates to refrigerant compositions comprising the present azeotrope-like compositions.

The refrigerant compositions of the present invention may be used in any of a wide variety of refrigeration systems including air-conditioning, refrigeration, heat-pump, HVAC systems, and the like. In certain preferred embodiments, the compositions of the present invention are used in refrigeration systems originally designed for use with an HFC refrigerant, such as, for example, HFC-134a. The preferred compositions of the present invention tend to exhibit many of the desirable characteristics of HFC-134a and other HFC refrigerants, including a GWP that is as low, or lower than that of conventional HFC refrigerants and a capacity that is as high or higher than such refrigerants. In addition, the relatively constant boiling nature of the compositions of the present invention makes them even more desirable than certain conventional HFCs for use as refrigerants in many applications.

In certain other preferred embodiments, the present compositions are used in refrigeration systems originally designed for use with a CFC-refrigerant. Preferred refrigeration compositions of the present invention may be used in refrigeration systems containing a lubricant used conventionally with CFC-refrigerants, such as mineral oils, silicone oils, polyalkylene glycol oils, and the like, or may be used with other lubricants traditionally used with HFC refrigerants. As used herein the term "refrigeration system" refers generally to any system or apparatus, or any part or portion of such a system or apparatus, which employs a refrigerant to provide cooling. Such refrigeration systems include, for example, air conditioners, electric refrigerators, chillers, transport refrigeration systems, commercial refrigeration systems and the like.

Any of a wide range of methods for introducing the present refrigerant compositions to a refrigeration system can be used in the present invention. For example, one method comprises attaching a refrigerant container to the low-pressure side of a refrigeration system and turning on the refrigeration system compressor to pull the refrigerant into the system. In such embodiments, the refrigerant container may be placed on a scale such that the amount of refrigerant composition entering the system can be monitored. When a desired amount of refrigerant composition has been introduced into the system, charging is stopped. Alternatively, a wide range of charging tools, known to those of skill in the art, is commercially available. Accordingly, in light of the above disclosure, those of skill in the art will be readily able to introduce the refrigerant compositions of the present invention into refrigeration systems according to the present invention without undue experimentation.

According to certain other embodiments, the present invention provides refrigeration systems comprising a refrigerant of the present invention and methods of producing heating or cooling by condensing and/or evaporating a composition of the present invention. In certain preferred embodiments, the methods for cooling an article according to the present invention comprise condensing a refrigerant composition comprising an azeotrope-like composition of the present invention and thereafter evaporating said refrigerant composition in the vicinity of the article to be cooled. Certain preferred methods for heating an article comprise condensing a refrigerant composition comprising an azeotrope-like composition of the present invention in the vicinity of the article to be heated and thereafter evaporating said refrigerant composition. In light of the disclosure herein, those of skill in the art will be readily able to heat and cool articles according to the present inventions without undue experimentation.

In another embodiment, the azeotrope-like compositions of this invention may be used as propellants in sprayable compositions, either alone or in combination with known propellants. The propellant composition comprises, more preferably consists essentially of, and, even more preferably, consists of the azeotrope-like compositions of the invention. The active ingredient to be sprayed together with inert ingredients, solvents, and other materials may also be present in the sprayable mixture. Preferably, the sprayable composition is an aerosol. Suitable active materials to be sprayed include, without limitation, cosmetic materials such as deodorants, perfumes, hair sprays, cleansers, and polishing agents as well as medicinal materials such as anti-asthma and anti-halitosis medications.

Yet another embodiment of the present invention relates to a blowing agent comprising one or more azeotrope-like compositions of the invention. In other embodiments,

the invention provides foamable compositions, and preferably polyurethane and polyisocyanurate foam compositions, and methods of preparing foams. In such foam embodiments, one or more of the present azeotrope-like compositions are included as a blowing agent in a foamable composition, which composition preferably includes one or more additional components capable of reacting and foaming under the proper conditions to form a foam or cellular structure, as is well known in the art. Any of the methods well known in the art, such as those described in "Polyurethanes Chemistry and Technology," Volumes I and II, Saunders and Frisch, 1962, John Wiley and Sons, New York, NY, which is incorporated herein by reference, may be used or adapted for use in accordance with the foam embodiments of the present invention.

Other uses of the present azeotrope-like compositions include use as solvents, cleaning agents, and the like. Those of skill in the art will be readily able to adapt the present compositions for use in such applications without undue experimentation.

**EXAMPLE**

The invention is further illustrated in the following example which is intended to be illustrative, but not limiting in any manner.

5    Example 1

An ebulliometer consisting of vacuum jacketed tube with a condenser on top which is further equipped with a Quartz Thermometer K96S4771 is used. About 35 g CF<sub>3</sub>I is charged to the ebulliometer and then HFO-1234ze is added in small, measured increments.

10    Temperature depression is observed when HFO-1234ze is added to CF<sub>3</sub>I, indicating a binary minimum boiling azeotrope is formed. From greater than about 0 to about 43 weight percent HFO-1234ze, the boiling point of the composition changed by about 2°C or less. The binary mixtures shown in Table 1 were studied and the boiling point of the compositions changed by about 2°C or less. The compositions exhibit azeotrope and/or azeotrope-like properties over this range.

15

Table 1HFO-1234/CF<sub>3</sub>I compositions at 14.42 psia

Wt.% HFO-1234	Wt.% CF <sub>3</sub> I	Temperature (°C)
0.00	100.00	-22.008
0.56	99.44	-22.085
5.13	94.87	-23.344
9.27	90.73	-23.730
14.26	85.74	-24.175
17.95	82.05	-24.300
23.12	76.88	-24.377
27.55	72.45	-24.339
32.11	67.89	-24.184
35.93	64.07	-24.020
38.96	61.04	-23.875
39.18	60.82	-23.836
42.43	57.57	-23.740